

Success	PI/ Award Year	Publication
<b>Cancer Treatments</b>		
<i>Treatment options for epithelial cancers by developing strategies for therapeutic immune radiotherapy delivering radioactivity directed specifically to tumors and tumor cells</i>	Pastan, I. 1999	Pai-Scherf, Lee H., et al. "Imaging and phase I study of 111In- and 90Y-labeled anti-LewisY monoclonal antibody B3." <i>Clinical Cancer Research</i> 6.5 (2000): 1720-1730
<i>Small lymphocytic lymphoma (SLL)/Chronic lymphocytic leukemia (CLL) therapy by developing approaches of anti-CD20 mAbs that preserve CD20 cell-surface expression on CLL cells which have led to new Rituximab-based combinations, novel schedules, and the design of the next generation of antibodies</i>	Weistner, A. 2007	Aue, Georg, et al. "Fractionated subcutaneous rituximab is well-tolerated and preserves CD20 expression on tumor cells in patients with chronic lymphocytic leukemia." <i>Haematologica</i> 95.2 (2010): 329-332
<b>AIDS Treatments</b>		
<i>Eliminating HIV-1 reservoirs using combination of intense treatment and immune intervention to eliminate activated latently infected cells</i>	Maldarelli, F. 2008	Maldarelli, Frank. "Targeting viral reservoirs: ability of antiretroviral therapy to stop viral replication." <i>Current Opinion in HIV and AIDS</i> 6.1 (2011): 49-56
<i>Understanding the interactions between hemolysis and infectious vasculopathy which modified the prevalence, age of onset, and severity of pulmonary hypertension among sickle cell patients</i>	Ognibene, F. 2001	Gladwin, Mark T., et al. "Pulmonary hypertension as a risk factor for death in patients with sickle cell disease." <i>New England Journal of Medicine</i> 350.9 (2004): 886-895
<b>Rare Diseases</b>		
<i>BtB support enabled identification of biomarkers (patent #US 20090286272 A1) and laid the groundwork for a Phase I clinical trial to evaluate the safety and effectiveness of cyclodextrin as a potential therapy for Niemann-Pick type C1. Additionally, the collaboration established by two BtB awards led the team to successfully apply for a U01 award, "A Phase 1 Dose Escalation Study of Vorinostat in Niemann-Pick C1 Disease"</i>	Porter, F. 2010 & 2011	Cluzeau, Celine VM, et al. "Microarray expression analysis and identification of serum biomarkers for Niemann-Pick disease, type C1." <i>Human Molecular Genetics</i> (2012): dds193
<b>Signaling Molecules and Genetic Disorders</b>		
<i>Lysosomal storage Disorder and Infantile Neuronal ceroid lipofuscinoses (INCLs) studies resulted in identifying Cysteamine Bitartrate (Cystagon) as a potential target that could stop the progression of retinal and brain damage in infants with INCL.</i>	Mukherjee, A. 2000.	Miao, Ning, et al. "Children with infantile neuronal ceroid lipofuscinosis have an increased risk of hypothermia and bradycardia during anesthesia." <i>Anesthesia and Analgesia</i> 109.2 (2009): 372
<i>Hutchinson-Gilford Progeria Syndrome (HGPS) and the use of Farnesyl Transferase inhibitors was investigated and showed a potential approach to tackle this disorder's effects on body weight, fat tissue and bone mineralization, and fracture rate.</i>	Gordon, L. 2005	Merideth, Melissa A., et al. "Phenotype and course of Hutchinson-Gilford progeria syndrome." <i>New England Journal of Medicine</i> 358.6 (2008): 592-604